Note :

“The Aprotinin content in Tisseel is purely synthetic in origin and there is no bovine content in Tisseel. The Tisseel Product Monograph will be revised shortly to reflect this change and will be posted in baxter.ca”

PRODUCT MONOGRAPH

Pr TISSEEL

Fibrin Sealant (Human), Vapor Heated, Solvent/Detergent Treated
500 IU (Fast Set)

in two presentation forms:
- Kit: Lyophilized Powders with Diluents for Topical Application,

- Frozen: Solutions for Thawing for Topical Application,

Hemostatic Agent

Manufactured by:	**Imported and Distributed by:**	**Date of Revision:**
Baxter AG A-1220 Vienna Austria	Baxter Corporation Mississauga, ON Canada L5N 0C2	March 05, 2013

Submission Control No: 160973 Date of Approval: March 21, 2013

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TISSEEL

Fibrin Sealant (Human), Vapor Heated, Solvent/Detergent Treated, 500 IU/mL

1. PART I: HEALTH PROFESSIONAL INFORMATION

1.1 SUMMARY PRODUCT INFORMATION

Route of Administration	Dosage Form / Strength	Clinically Relevant Nonmedicinal Ingredients
Topical	**500 IU (Fast Set) - TISSEEL** -Lyophilized Powders with Diluents: 2.0 mL, 4.0 mL, 10.0 mL (total volume) or - Frozen Solutions for Thawing: 2.0 mL, 4.0 mL, 10.0 mL (total volume)	*For a complete listing see Dosage Forms, Composition and Packaging Section.*

1.2 DESCRIPTION

TISSEEL (Fibrin Sealant (Human), Vapor Heated, Solvent/Detergent Treated) is a two-component fibrin sealant made from pooled human plasma. When combined, the two components, Sealer Protein (Human) and Thrombin (Human), mimic the final stage of the blood coagulation cascade. TISSEEL is intended only for topical administration.

Sealer Protein (Human)

Sealer Protein (Human) is a sterile, non-pyrogenic, vapor-heated and solvent/detergent treated preparation made from pooled human plasma. Sealer Protein (Human) is provided either as a freeze-dried powder Sealer Protein Concentrate (Human) for reconstitution with Aprotinin Solution (Bovine or Synthetic) or as a finished frozen liquid Sealer Protein-Aprotinin Solution pre-filled into one side of a double-chamber syringe (chamber containing Sealer Protein - Aprotinin Solution is marked as "1").

The active ingredient in Sealer Protein (Human) component is fibrinogen. In addition Factor XIII is co-purified with clottable protein from human plasma. No Factor XIII is added to the Sealer Protein (Human) manufacturing process, resulting in a Factor XIII level of 0.6-10 U/mL in the drug product. A Fibrinolysis Inhibitor, Aprotinin (Bovine or Synthetic) is included in the Sealer Protein (Human) component to preclude premature fibrinolysis. To obtain Sealer Protein

Concentrate (Human), cryoprecipitate derived from the plasma is dissolved in buffer solution, solvent/detergent treated, purified by precipitation and washing steps, vapor heat treated, formulated, sterile filtered and either freeze-dried in vials or concentrated under vacuum and frozen in pre-filled syringes.

Thrombin (Human)

Thrombin (Human) is a sterile, non-pyrogenic, vapor-heated and solvent/detergent treated preparation made from pooled human plasma. Thrombin (Human) is also provided either as a freeze-dried powder for reconstitution with Calcium Chloride Solution or as a finished frozen liquid Thrombin-Calcium Chloride Solution pre-filled into one side of a double-chamber syringe (chamber containing Thrombin-Calcium Chloride solution is marked as "2").

The active ingredient human Thrombin is prepared from plasma through a series of separation and filtration steps followed by incubation of the solution with calcium chloride to activate prothrombin to thrombin. The solution subsequently undergoes ultra/diafiltration, vapor heat treatment, solvent/detergent treatment, purification by ion exchange chromatography, formulation, sterile filtration and either freeze-drying in vials or filling and freezing in pre-filled syringes.

The setting rate of the Fibrin Sealant depends on the concentration of the human Thrombin as contained in the Thrombin-Calcium Chloride Solution used. The TISSEEL thrombin component is provided in a 500 IU/mL concentration (TISSEEL 500 IU, Fast Set). Time to clot formation/polymerisation will be completed within seconds with a Thrombin concentration of 500 IU/mL. Therefore, TISSEEL 500 IU (Fast Set) is indicated to be used to achieve rapid setting, e.g. for hemostasis.

Sealer Protein (Human) and Thrombin (Human) are made from pooled human plasma. The vapor heat and solvent/detergent treatment steps used in the manufacturing process have been shown to be capable of significant viral reduction. Despite this, when medicinal products prepared from human blood or plasma are administered, the possibility of transmitting infective agents cannot be totally excluded *(see WARNINGS AND PRECAUTIONS and PHARMACEUTICAL INFORMATION, and* see *DOSAGE FORMS, COMPOSITION AND PACKAGING* for available package sizes and presentations).

1.3 INDICATIONS AND CLINICAL USE

TISSEEL (Fibrin Sealant (Human), Vapor Heated, Solvent Detergent Treated) is used in addition to standard measures, to achieve hemostasis, to seal or glue tissue, and to support wound healing.

Indications include:
abdominal surgery[2, 6, 7]
cardiovascular surgery[4, 5, 8, 9]
orthopedic surgery[1, 10]
thoracic surgery[11, 12, 13]
urology[3, 14, 15, 16]

Geriatrics (> 65 years of age):
Efficacy and safety in use of Fibrin Sealant has been evaluated in a clinical trial involving a number of geriatric patients. No overall differences in safety or effectiveness were observed between these subjects and younger subjects.

Pediatrics (1.1 – 16 years of age) or (< 16 years of age):
Efficacy and safety in use of Fibrin Sealant has been evaluated in a clinical trial involving also a group of pediatric patients and was not found to be different from an adult population.

1.4 CONTRAINDICATIONS

- Patients who are hypersensitive to this drug or to any ingredient in the formulation or component of the container. For a complete listing, see the *DOSAGE FORMS, COMPOSITION AND PACKAGING* Section of the product monograph.
  - For TISSEEL (Fibrin Sealant (Human), Vapor Heated, Solvent/Detergent Treated) containing Aprotinin of bovine or synthetic source: Known hypersensitivity to bovine protein.
  - For TISSEEL (Fibrin Sealant (Human), Vapor Heated, Solvent/Detergent Treated) generally: TISSEEL should not be used in individuals with a known hypersensitivity to Aprotinin.
- TISSEEL is contraindicated for intravascular application. Intravascular application may result in life threatening thromboembolic events.
- For the treatment of massive and brisk arterial or venous bleeding, TISSEEL alone is not effective and not indicated for this treatment.

1.5 WARNINGS AND PRECAUTIONS

Serious Warnings and Precautions

Intravascular application of TISSEEL can lead to intravascular coagulation, may result in life-threatening thromboembolic events and might increase the likelihood and severity of acute hypersensitivity reactions in susceptible patients. Especially in coronary bypass surgery, TISSEEL should be applied with caution to minimize any risk of intravascular application.

However, if in well-founded cases the injection of TISSEEL and/or Thrombin Solution/s into a tissue or vessel is indicated, careful risk/benefit analysis of the individual case is to be carried out.

As with any other protein products, hypersensitivity or allergic/anaphylactic/anaphylactoid reactions may occur with the use of TISSEEL, a fibrin sealant, that contains aprotinin. In isolated cases, these reactions have progressed to severe anaphylaxis. Aprotinin, a monomeric polypeptide, is known to be associated with anaphylactic reactions. These reactions may also occur in patients receiving Aprotinin or TISSEEL for the first time or even if first application was well tolerated, but generally risk may be increased if the preparation is applied repeatedly over time or in the same setting. Manifestations of hypersensitivity reactions to TISSEEL observed include: flush, urticaria, pruritus, nausea, drop in blood pressure, tachycardia or bradycardia, dyspnea, severe hypotension, bronchospasm, wheezing, angioedema, erythema, paresthesia. Fatal anaphylactic reactions, including anaphylactic shock, have also been reported with TISSEEL (see ADVERSE REACTIONS).

In the event of hypersensitivity reactions, administration of TISSEEL is to be discontinued. Remove any applied, polymerized product from the surgical site. Mild reactions can be managed with antihistamines. Severe hypotensive reactions require immediate intervention using current principles of shock therapy. Adequate medical treatment and provisions should be available for immediate use in the event of an anaphylactic reaction.

Caution must be used when applying fibrin sealant using pressurized air or gas (See DOSAGE AND ADMINISTRATION). Life-threatening/fatal air or gas embolism has occurred with the use of spray devices employing pressure regulator to administer fibrin sealants. This event appears to be related to the use of the spray device at higher than recommended pressures and in close proximity to the tissue surface. The risk appears to be higher when fibrin sealants are sprayed with air, as compared to CO_2, when TISSEEL is used in open wound surgery.

Solutions containing alcohol, iodine or heavy metals will interfere with the product's performance due to denaturation of proteins or other mechanisms.

General

Fibrin Sealant is made from human plasma. Products made from human plasma may contain infectious agents, such as viruses, that can cause disease. This also applies to unknown or emerging viruses and other pathogens. The risk that such products will transmit an infectious agent has been reduced by screening plasma and by inactivating and removing certain viruses. Despite these measures, there may still carry a risk of transmitting infectious agents, e.g., viruses, and theoretically, the Creutzfeldt-Jakob disease (CJD) agent.

The standard measures taken (including double virus inactivation by vapor heat treatment and solvent detergent treatment) are considered effective for enveloped viruses such as HIV, HBV, and HCV, and for the non-enveloped virus HAV.

Some viruses, such as parvovirus B19, are particularly difficult to remove or inactivate at this time. Parvovirus B19 most seriously affects pregnant women (fetal infection) and individuals with immunodeficiency or increased red blood cell turnover.

In the submucous injection of fibrin sealant into hollow organs to stop GI-bleeding (stomach, duodenum), the following points are to be considered:

- Insertion of the needle into the organ wall may result in accidental perforation, which in rare cases may injure adjoining organs or vessels.

- Injection into the submucous membrane may cause a mechanical dissection between the tunica mucosa and the tunica muscularis propria, which in rare cases may lead to vessel injury or the formation of an intramural hematoma.

- Injection into the nasal mucosa must be avoided, as severe allergic-anaphylactoid reactions have been seen and thromboembolic events may occur.

This product must not be used in animals.

The user is cautioned against the spray application of TISSEEL with devices produced by other manufacturers. Only use application devices licensed for the administration of TISSEEL.

TISSEEL with the Spray Set must not be used in enclosed body areas.

When spraying TISSEEL, changes in blood pressure, pulse, oxygen saturation and end tidal CO_2, should be monitored because of the possibility of occurrence of air or gas embolism.

The EASYSPRAY control device, DuploSpray MIS Surgical Sealant Delivery System and the Spray Set may be obtained from Baxter.

To prevent TISSEEL from adhering to gloves and instruments, wet these with saline before contact with Sealant.

To avoid the formation of excess granulation tissue and to ensure gradual absorption of the solidified fibrin sealant, only a thin layer of TISSEEL should be applied (see Application Precautions and *OVERDOSAGE*).

Immediately before application, expel and discard the first several drops from the application cannula to ensure adequate mixing of the Sealer Protein and Thrombin Solutions.

Application Precautions

Only a thin layer of TISSEEL should be applied. Excessive thickness of the fibrin sealant layer may interfere with the products efficacy and the wound healing process.
It is strongly recommended that every time TISSEEL is applied to a patient, the name and batch number of the product are recorded in order to maintain a link between the patient and the batch of the product.

When applying TISSEEL using the EASYSPRAY device, be sure to use the pressure within the pressure range recommended by the spray device manufacturer. In the absence of a specific recommendation avoid using pressure above 20-25 psi. Do not spray closer than the distance recommended by the spray device manufacturer. In the absence of a specific recommendation avoid spraying closer than 10-15 cm from the surface of the tissue. When spraying TISSEEL, changes in blood pressure, pulse, oxygen saturation and end tidal CO_2 should be monitored because of the possibility of occurrence of air or gas embolism.

When applying TISSEEL using the DuploSpray MIS system connect to regulated CO_2, gas source only; maximum input pressure not to exceed 100 psi. To avoid possible gas embolism, do not directly spray into circulatory pathways. Any application of pressurized gas may be associated with a potential risk of gas embolus, tissue rupture or gas entrapment with compression, which may be life threatening. Be sure to take appropriate measures to address these risks by observing the recommended minimum spraying distance of 2 cm (optional working distance 3cm) and the maximal flow rate of 2.0 standard liters per minute. For use by or on order of a licensed physician.

When applying TISSEEL using a spray device be sure to use a pressure and a distance from tissue with the ranges recommended by the manufacturer as follows:

<table>
<tr><td colspan="6">Recommended pressure, distance and devices for spray application of TISSEEL</td></tr>
<tr><td>Surgery</td><td>Spray set to be used</td><td>Applicator tips to be used</td><td>Pressure regulator to be used</td><td>Recommended distance from target tissue</td><td>Recommended Spray Pressure</td></tr>
<tr><td rowspan="2">Open wound</td><td>Tisseel/Artiss Spray Set</td><td>n.a</td><td>EasySpray</td><td rowspan="2">10-15cm</td><td rowspan="2">1.5-2.0 bar (21.5-28.5 psi)</td></tr>
<tr><td>Tisseel/Artiss Spray set 10 pack</td><td>n.a</td><td>EasySpray</td></tr>
<tr><td colspan="6"></td></tr>
<tr><td rowspan="8">Laparoscopic/ minimally invasive procedures</td><td rowspan="8">n.a</td><td rowspan="2">Duplospray MIS Applicator 20cm</td><td>Duplospray MIS Regulator</td><td rowspan="8">2-5cm</td><td rowspan="8">1.2-1.5bar (18-22 psi)</td></tr>
<tr><td>Duplospray MIS Regulator NIST B11</td></tr>
<tr><td rowspan="2">Duplospray MIS Applicator 30cm</td><td>Duplospray MIS Regulator</td></tr>
<tr><td>Duplospray MIS Regulator NIST B11</td></tr>
<tr><td rowspan="2">Duplospray MIS Applicator 40cm</td><td>Duplospray MIS Regulator</td></tr>
<tr><td>Duplospray MIS Regulator NIST B11</td></tr>
<tr><td rowspan="2">Replaceable Tip</td><td>Duplospray MIS Regulator</td></tr>
<tr><td>Duplospray MIS Regulator NIST B11</td></tr>
</table>

Solutions containing alcohol, iodine or heavy metals will interfere with the product's performance due to denaturation of proteins or other mechanisms. If any of these substances have been used to clean the wound area, the area must be thoroughly rinsed before application of TISSEEL.

Cardiovascular
In coronary bypass surgery, TISSEEL should be applied with caution to minimize any risk of intravascular application.

Sexual Function/Reproduction
Animal reproduction studies have not been conducted with TISSEEL. It is also not known whether it can affect reproduction capacity.

Special Populations

Pregnant Women: No experience.
Animal reproduction studies have not been conducted with TISSEEL. It is also not known whether it can cause fetal harm when administered to a pregnant woman. Healthcare providers should balance the potential risks and only prescribe TISSEEL if clearly needed.

See WARNING AND PRECAUTIONS for information on Parvovirus B19 infection.

Nursing Women:
It is not known whether this drug is excreted in human milk. Because many drugs are excreted in human milk, caution should be exercised when TISSEEL is administered to a lactating woman. Healthcare providers should balance the potential risks and only prescribe TISSEEL if clearly needed.

Neurosurgical Procedures

The safety and effectiveness of TISSEEL used alone or in combination with biocompatible carriers in neurosurgical procedures or other surgeries involving confined spaces have not been evaluated. If fibrin sealants are applied in confined spaces, the risk of compressive complications should be taken into account.

1.6 ADVERSE REACTIONS

Adverse Drug Reaction Overview

Allergic and/or anaphylactic reactions may occur in patients with a history of hypersensitivity against bovine protein and/or Aprotinin. Such reactions may be seen in the event of repeated administration, even if the first application was well tolerated. However, allergic and/or anaphylactic reactions may also occur in patients receiving TISSEEL for the first time.

If symptoms require treatment to be initiated, this should be effected in the usual manner, as for instance with antihistamines, corticoids or adrenalin. No adverse events of this type were reported during the clinical trials for TISSEEL.

Clinical Trial Adverse Drug Reactions (ADRs)

Because clinical trials are conducted under widely varying conditions, adverse reaction rates observed in the clinical trials may not reflect the rates observed in practice.

TISSEEL 500 IU (Fast Set):

The following ADRs have been reported from three clinical trials. None of the events was classified as serious.

During a clinical study in cardiovascular surgery where 157 patients were treated with TISSEEL 500 IU (Fast Set) two cases of increased Fibrin D-Dimer levels were observed, but did not exceed values reported in the literature occurring after this type of surgery. Postoperatively increased D-Dimers may result at least partly from the degradation of Fibrin Sealant (*see Part II Clinical Trials, trial number 550003*).

In a study in axillary lymph node dissection where 79 patients were treated with TISSEEL 500 IU (Fast Set), the most frequently occurring ADRs were seroma (13 cases) and pain in limb (4 cases). However, the overall incidence of (related and unrelated) seroma and procedural pain was comparable between the TISSEEL 500 IU (Fast Set) and an untreated control group (21.5% vs. 18.3%) in this study. Other adverse events reported were: 2 cases each of nausea, post-operative wound infection and post-procedural pain, and 1 case each of pain, increase of body temperature, increase of fibrin degradation products, sensory disturbance, rash, axillary vein thrombosis and hypotension.

No adverse events related to TISSEEL 500 IU (Fast Set) were reported from a study in hip joint replacement from the 53 patients treated with the product.

The related adverse events of the cardiovascular surgery study and the lymph node dissection study are listed according to their system organ class in the table below. Frequencies are based on patients treated in the respective studies because the indications of the two studies were too different to pool the data.

System Organ Class	Adverse events (Preferred Term)	TISSEEL 500 IU Number of events/Number of patients treated
Blood and lymphatic system disorders	Fibrin D-Dimer increased	2 of 157
	Increase of Fibrin degradation products	1 of 79
Nervous system disorders	Sensory disturbance	1 of 79
Gastrointestinal disorders	Nausea	2 of 79
Skin and subcutaneous tissue disorders	Rash	1 of 79
Musculosceletal and connective tissue disorders	Pain in extremity	4 of 79
Infections and infestations	Postoperative wound infection	2 of 79
Injury, poisoning and procedural complications	Seroma	13 of 79
Vascular disorders	Axillary vein thrombosis	1 of 79
	Hypotension	1 of 79
General disorders and administration site conditions	Procedural Pain	2 of 79
	Pain	1 of 79
	Body temperature increased	1 of 79

Post-Market Adverse Drug Reactions

The adverse reactions listed below reflect what has been reported in post-marketing experience with Baxter Fibrin Sealants:

Immune system disorders:

Anaphylactic responses, hypersensitivity (manifestations include bradycardia, tachycardia, hypotension, flushing, bronchospasm, wheezing, dyspnea, nausea, urticaria, angioedema, pruritus, erythema, paresthesia; anaphylactic reactions, including anaphylactic shock, have included fatal outcomes)

Cardiac disorders:
Bradycardia, tachycardia

Vascular disorders:
Hypotension, haematoma, embolism arterial, including cerebral artery embolism, cerebral infarction*

*as a result of intravascular application into the superior petrosal sinus

Respiratory, thoracic and mediastinal disorders:
Dyspnoea

Gastrointestinal disorders:
Nausea, intestinal obstruction

Skin and subcutaneous tissue disorders:
Urticaria

General disorders and administration site conditions:
Flushing, impaired healing, oedema, pyrexia

Injury, poisoning and procedural complication:
Seroma

There have been rare reports of serious adverse events such as paralysis and other compressive complications possibly related to the use of fibrin sealant in combination with resorbable hemostatic agents.

Other adverse reactions associated with products of the fibrin sealant/hemostatic class include: air or gas embolism when using devices with pressurized air or gas; this event appears to be related to the use of the spray device at higher than recommended pressures and in close proximity to the tissue surface. Manifestations of hypersensitivity or allergic reactions include application site irritation, chest discomfort, chills, headache, lethargy, restlessness, and vomiting.

There have also been rare reports of fatalities following the misadministration of topical thrombin.

There have been rare reports of air embolism associated with misapplication of fibrin sealant using a spray device.

Because these reactions are reported voluntarily and the population is of uncertain size, it is not always possible to reliably estimate the frequency of these reactions.

1.7 DRUG INTERACTIONS

Overview

No formal interaction studies have been performed.[1] Drug Interactions are not known. TISSEEL (Fibrin Sealant (Human), Vapor Heated, Solvent/Detergent Treated) can even be

[1] None known based upon the absence of data from clinical trials, current medical/scientific literature, and safety reports.

applied in fully heparinised patients (e.g. extracorporeal circulation).

Incompatibilities

Solutions containing alcohol, iodine or heavy metals will interfere with the product's performance due to denaturation of proteins or other mechanisms. If any of these substances have been used to clean the wound area, the area must be thoroughly rinsed before application of TISSEEL.

Oxidized cellulose-containing preparations may reduce the efficacy of TISSEEL by inactivating thrombin and should not be used as carrier materials.

TISSEEL must not be mixed with other medicinal products.

1.8 DOSAGE AND ADMINISTRATION

TISSEEL 500 IU (Fast Set) (Fibrin Sealant (Human), Vapor Heated, Solvent/Detergent Treated) is available in two different dosage forms: Kit and Frozen.

TISSEEL Kit consists of 2 vials containing powder preparations of Sealer Protein Concentrate and Thrombin and two vials containing liquid preparations of Aprotinin and Calcium Chloride. The Sealer Protein Concentrate is reconstituted with the Aprotinin and produces a solution that is referred to as the Sealer Protein-Aprotinin Solution. Thrombin is reconstituted with the Calcium Chloride Solution, which produces a solution that is referred to as the Thrombin-Calcium Chloride Solution. These two solutions are applied using an application device described in *DOSAGE AND ADMINISTRATION, Method of Application Section*. TISSEEL Kit is available in 2, 4 and 10 mL pack sizes (1, 2 and 5 mL reconstituted Sealer Protein Solution and 1, 2 and 5 mL reconstituted Thrombin Solution).

TISSEEL Frozen consists of a pre-filled double-chamber syringe containing Sealer Protein-Aprotinin Solution and Thrombin-Calcium Chloride Solution. TISSEEL Frozen is available in 2, 4 and 10 mL pack sizes.

Dosing Considerations

For topical use only – do not inject.
Prior to applying TISSEEL the surface area of the wound needs to be dried by standard techniques (e.g. intermittent application of compresses, swabs, use of suction devices)

Recommended Dose and Dosage Adjustment

Dosage of TISSEEL, i.e., the amount applied, is individualized based on the application area and purpose of use. Only a thin layer of TISSEEL should be applied. Excessive thickness of the fibrin layer may interfere with the product's efficacy and the wound healing process.

The application can be repeated, if necessary. However, avoid re-application to a pre-existing polymerized TISSEEL layer, as TISSEEL may not adhere firmly to a polymerized layer.

If used for tissue adherence, it is recommended that the initial application cover the entire intended application area.

The amount of TISSEEL to be applied depends on the size of the surface to be covered. The approximate surface areas covered by each package size of TISSEEL are listed in the following table:

Area to be sealed (cannula, catheter)	**Area to be sealed using compressed gas (spray application)**	**Required package size of TISSEEL**
8 cm^2	25-100 cm^2	2 mL
16 cm^2	50-200 cm^2	4 mL
40 cm^2	125-500 cm^2	10 mL

Administration

Various methods can be used to simultaneously apply the two components of TISSEEL:

a) using application cannula contained in DUPLOJECT System / DUO Set
b) using Spray Set and EASYSPRAY
c) using DUPLOCATH Application Catheters or other accessories provided by Baxter
d) using DUPLOSPRAY MIS regulator, applicator and tips

1. Preparation of TISSEEL Kit (Freeze-Dried)

The following instructions are for the reconstitution of TISSEEL Kit using the DUPLOJECT System provided within the Kit. However, an equivalent device can also be used. For operation instructions please refer to the Instructions for Use provided together with this device.

Check for appropriate component colour-coding before reconstitution of the component.

Prior to reconstitution of TISSEEL the rubber stoppers of all vials should be cleansed.

After reconstitution, the product must be used within 4 hours.

Reconstituted solutions must not be refrigerated or frozen. Vials are for single use only.

Note: Do not use the syringes and needles previously used for reconstitution of one component for the reconstitution of the other component to prevent premature setting.

Direct contact between disinfectant and product must be avoided (See *DRUG INTERACTIONS*).

a) How to Prepare TISSEEL Sealer Protein Solution

Freeze-dried Sealer Protein Concentrate is reconstituted in the Aprotinin Solution of 3,000 KIU/mL using the FIBRINOTHERM warming and stirring device (recommended method). Alternatively, a sterile water bath at a temperature of 33°C to 37°C can be used.

i) Reconstitution of Freeze-Dried Sealer Protein Concentrate Using FIBRINOTHERM device:

For ease of handling, a combined heating and stirring device, FIBRINOTHERM, has been developed to meet the specific requirements of reconstituting freeze-dried Sealer Protein Concentrate. FIBRINOTHERM is a thermoblock with a magnetic stirrer (the vials for freeze-dried Sealer Protein Concentrate contain a magnetic spin propeller to stir the contents). Heating and stirring can be operated independently. In a first step, FIBRINOTHERM heats up to 37°C and then maintains that temperature constantly with minimum variation. The signal light will turn on and off as FIBRINOTHERM is maintaining its temperature. FIBRINOTHERM has been designed to hold the various vial sizes of freeze-dried Sealer Protein Concentrate and Aprotinin Solution.

- After plugging the FIBRINOTHERM into an electrical socket and activating the warmer using the amber switch, place the Sealer Protein Concentrate and Aprotinin Solution vials from the TISSEEL Kit into prewarmed warming wells with the appropriate sized adaptor ring. Refrigerated product may take up to 5 minutes to warm. Room temperature product will take less time.
- Transfer Aprotinin Solution into vial containing freeze-dried Sealer Protein Concentrate using blue-scaled syringe of corresponding size (or syringe that has been used for dilution of Aprotinin Solution).
  Note: Only combine preheated Aprotinin Solution with preheated Sealer Protein Concentrate.
- Place vial into largest opening of FIBRINOTHERM (if necessary, use adaptors). Turn on stirrer with flip switch and stir contents for 8 – 10 minutes.
- Reconstitution of freeze-dried Sealer Protein Concentrate is complete as soon as no undissolved particles are detectable in transparent light. Otherwise, replace into FIBRINOTHERM and agitate for another few minutes until the solution appears homogeneous.
  Note: Excessive stirring (20 minutes or more) may compromise product quality.

Note: Keep the Sealer Protein Solution at 37°C or at room temperature without stirring if it is not to be used immediately. Before use the solution should be warmed to 37°C. To ensure homogeneity, stir or swirl briefly before aseptically drawing up the Sealer Protein Solution into the blue-scaled syringe provided in DUPLOJECT System.

ii) Reconstitution of Freeze-Dried Sealer Protein Concentrate Using a Water Bath:

Note: When using a water bath for reconstitution, special precautions must be taken against submerging the vial, particularly the septum, to avoid possible contamination.

- Preheat the vials containing the Sealer Protein Concentrate (lyophilized) and the Aprotinin Solution for approximately 3 minutes in a water bath at a temperature of 33°C to 37°C. (Heating beyond 37°C must be avoided!)
- Remove the flip-off caps from the vial containing freeze-dried Sealer Protein Concentrate and the vial containing Aprotinin Solution, disinfect the rubber stoppers of both vials with a germicidal solution and allow to dry. Do not use iodine-containing preparations such as betadine for disinfection.
- Transfer Aprotinin Solution into vial containing freeze-dried Sealer Protein Concentrate using blue-scaled syringe of corresponding size (or syringe that has been used for dilution of Aprotinin Solution).
- Allow vial to stand at 37°C for one minute.
- Swirl briefly and vigorously with a circular motion (avoid excessive frothing) and replace vial into water-bath for another 10 – 15 minutes.
- Reconstitution of freeze-dried Sealer Protein Concentrate is complete as soon as no undissolved particles are detectable in transparent light. Otherwise, swirl again briefly and keep vial at 37°C for a few more minutes.
- Draw up Sealer Protein Solution into a sterile blue-scaled syringe using aseptic precautions (insert a needle through the rubber stopper at its center to allow access of air).

Note: If not used immediately, keep Sealer Protein Solution at 37°C or at room temperature without stirring if it is not to be used immediately. Before use the solution should be warmed to 37°C. The maximum holding time of 4 hours upon reconstitution must not be exceeded. To ensure homogeneity, swirl with a circular motion (avoid frothing) before drawing up the solution.

b) How to Prepare Thrombin Solution

- Remove the flip-off caps from the vial containing Thrombin and the vial containing Calcium Chloride Solution, disinfect the rubber stoppers of both vials with a germicidal solution and allow to dry. Do not use iodine-containing preparations such as betadine for disinfection.
- Transfer the contents of the vial with Calcium Chloride Solution into the vial containing the freeze-dried Thrombin using the sterile reconstitution components provided with the DUPLOJECT System.
- Swirl briefly.
- Place the vial into the adapted opening of the FIBRINOTHERM device or Water Bath.

- Reconstitution of Thrombin is complete when all of the Thrombin concentrate is dissolved.
- Keep the Thrombin Solution at 37°C or at room temperature without stirring if it is not to be used immediately. Before use the solution should be warmed to 37°C.

c) Transferring to the Sterile Field

For transfer of the Sealer Protein Solution and the Thrombin Solution to the sterile field, the scrub nurse should withdraw the solutions while the circulating nurse holds the non-sterile vials. The solutions should be withdrawn slowly by firm constant aspiration to reduce the risk of large air bubbles.

2. Preparation of TISSEEL Frozen (in pre-filled syringes)

Do not use TISSEEL unless it is completely thawed and warmed (liquid consistency).

Do not remove the protective syringe cap until thawing is complete and application tip is ready to be attached.

Room Temperature Thawing

Approximate thawing times when using this method are:

Pack Size (In Pouches)	Thawing Time at Room Temperature
2 mL	60 minutes
4 mL	110 minutes
10 mL	160 minutes

Unopened pouches of TISSEEL 500 IU (Fast Set), thawed at room temperature, may be stored for up to 7 days at room temperature (15°C -25°C) after removal from the freezer. Prior to use, the product should be warmed to 33°C to 37°C.

TISSEEL in pre-filled syringe (frozen) may be thawed under controlled conditions using one of the three following options:

Option 1 – Thawing on the sterile field using a Sterile Water Bath:

33°C to 37°C sterile water bath - transfer double-chamber syringe set and the inner pouch to the sterile field, remove the double-chamber syringe with pre-filled syringes from inner pouch and place directly into sterile water bath. Ensure the double-chamber syringe is completely immersed under the water.

Approximate thawing and warming times when using this method are:

Pack Size	Thawing/Warming Times 33°C to 37°C Sterile Water Bath (Pouches Removed)
2 mL	5 minutes
4 mL	5 minutes
10 mL	12 minutes

Option 2 – Thawing off the sterile field using a Water Bath:

33°C to 37°C non-sterile water bath, double-chamber syringe in two pouches - maintain the double-chamber syringe set in both pouches and place into a water bath off the sterile field for appropriate time. Ensure the pouches remain submerged throughout thawing. Remove from the water bath after thawing, dry external pouch and transfer inner pouch and pre-filled syringes onto the sterile field.

Approximate thawing and warming times when using this method are:

Pack Size	Thawing/Warming Times 33°C to 37°C Non-Sterile Water Bath (In Pouches)
2 mL	30 minutes
4 mL	40 minutes
10 mL	80 minutes

Option 3 – Thawing off the sterile field using an Incubator:

Incubator (33°C to 37°C) in pouches – maintain the -pre-filled syringe in both pouches and place into an incubator for appropriate time. Remove from incubator after thawing and transfer inner pouch and pre-filled syringes onto the sterile field.

Approximate thawing and warming times when using this method are:

Pack Size	Thawing/Warming Times 33°C to 37°C Incubator (In Pouches)
2 mL	40 minutes
4 mL	85 minutes
10 mL	105 minutes

NOTE:
After thawing, the product must not be refrigerated, refrozen or be exposed to temperatures above 37^{o}C.

Keep the product at 33°C - 37^{o}C until needed.

The product must be used within 12 hours after warming to 33°C - 37°C or removal from original pouches.

3. Method of Application

Application Considerations

See *WARNINGS AND PRECAUTIONS*, General and Application Precautions.

The use of TISSEEL is restricted to experienced surgeons who have been trained in the use of TISSEEL.

In open wound surgery- a pressure regulator device that delivers a maximum pressure of no more than 2.0 bar (28.5 psi) should be used.

In minimally invasive/laparoscopic procedures- a pressure regulator device that delivers a maximum pressure of no more than 1.5 bar (22 psi) and uses carbon dioxide gas only should be used.

TISSEEL should only be reconstituted and administered according to the instructions and with the devices recommended for this product (see section 1.5)

Regarding spray applications, see section 1.5 for specific recommendations on the required pressure and distance from tissue per surgical procedure and length of applicator tips.

The setting rate depends on the concentration of the human Thrombin as contained in the Thrombin-Calcium Chloride solution used. The TISSEEL Thrombin component is provided in a 500 IU/mL concentration (TISSEEL 500 IU, Fast Set). A 4 IU/mL Thrombin concentration is also available and marketed as ARTISS 4IU, Slow Set.

Prior to application, TISSEEL must be warmed to 33°C to 37°C. TISSEEL must not be exposed to temperatures above 37°C and must not be microwaved.

The sealer protein and the thrombin solutions should be clear or slightly opalescent. Do not use solutions that are cloudy or have deposits. Inspect products for particulate matter and discoloration prior to application.

Before application of TISSEEL, ensure that parts of the body outside the desired application area are sufficiently covered to prevent tissue adhesion at undesired sites.

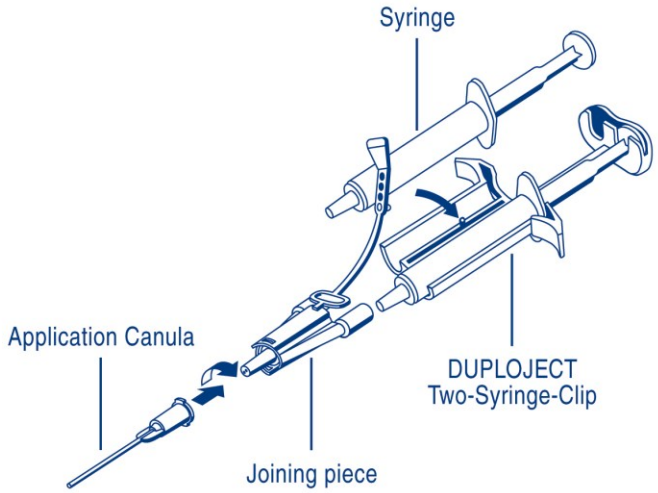
- Prior to applying TISSEEL the surface area of the wound needs to be dried by standard techniques (e.g. intermittent application of compresses, swabs, use of suction devices). Apply TISSEEL as a thin layer. The initial amount of the product to be applied should be sufficient to entirely cover the intended application area.

- If TISSEEL does not fully adhere to tissue and bleeding continues, remove TISSEEL clot and repeat application.
- Ensure that the two components are quickly and thoroughly mixed, which is essential for TISSEEL to gain the optimum strength.
- The Spray Set is particularly suitable for spraying of larger areas.
- In operation sites where access is difficult or when using an endoscope or trocar, TISSEEL can be applied using DUPLOCATH Application Catheters or DuploSpray MIS Surgical Sealant Delivery System.
- With the use of TISSEEL, 500 IU/mL, the clotting process will be completed within seconds.
- Once turbid, TISSEEL can no longer be manipulated.
- Solidified Sealant reaches its ultimate strength after about two hours (70% after about ten minutes).

a) Simultaneous Application using application cannula contained in DUPLOJECT System or DUO Set:

i) Administration using DUPLOJECT System

- For application, clip the two single-use syringes with the reconstituted Sealer Protein Solution and Thrombin Solution into the DUPLOJECT Two-Syringe Clip and connect this assembly to a joining piece and an application cannula.
- The common plunger of the DUPLOJECT Two-Syringe Clip ensures that equal volumes are fed through the joining piece, before being mixed in the application cannula and ejected.



Operating Instructions:

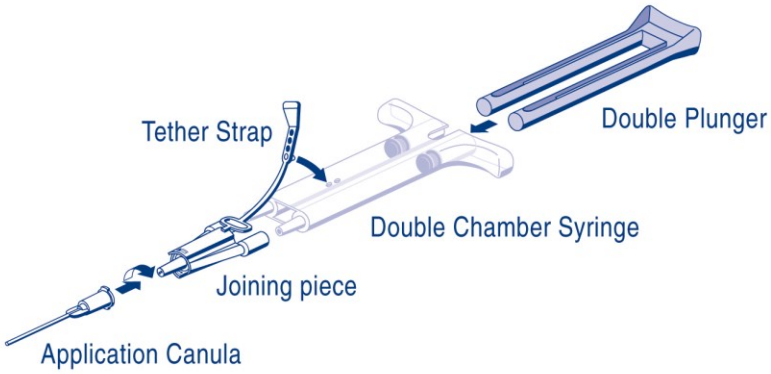
- Place the two syringes filled with Sealer Protein Solution and with Thrombin Solution into the clip. Both syringes should be filled with equal volumes.
- Connect the nozzles of the two syringes to the joining piece ensuring that they are firmly fixed. Secure the joining piece by fastening the tether strap to the DUPLOJECT Two-Syringe Clip. Should the pull strap tear, use the spare joining piece. If none is available, further use is still possible but tightness of the connection needs to be ensured to prevent any risk of leaking.
- Fit an application cannula onto the joining piece.
- Do not expel the air remaining inside the joining piece or application cannula until you start actual application as the aperture of the cannula may clog otherwise.
- Apply the mixed Sealer Protein - Thrombin Solution onto the recipient surface or surfaces of the parts to be sealed.

If application of the fibrin sealant components is interrupted, clogging may occur in the cannula. To resume application, replace the application cannula with a new one. If the apertures of the joining piece are clogged, use the spare joining piece provided in the DUPLOJECT System.

Note: For operation instructions please refer to the Instructions for Use provided together with the DUPLOJECT System.

ii) Administration using DUO Set

- For application, the double-chamber syringe with the Sealer Protein Solution and the Thrombin Solution has to be connected to a joining piece and an application cannula.



Operating Instructions:

- Remove the cap covering the nozzles of the double-chamber syringe.

- Connect the nozzles of the double-chamber syringe to the joining piece ensuring that they are firmly fixed. Secure the joining piece by fastening the tether strap to the double-chamber syringe. Should the pull strap tear, use the spare joining piece. If none is available, further use is still possible but tightness of the connection needs to be ensured to prevent any risk of leaking.
  Fit an application cannula onto the joining piece.
- Fit the double plunger to the end of the syringe chamber.
- Do not expel the air remaining inside the joining piece or application cannula until you start actual application as the aperture of the cannula may clog otherwise
- Apply the mixed Sealer Protein - Thrombin Solution onto the recipient surface or surfaces of the parts to be sealed

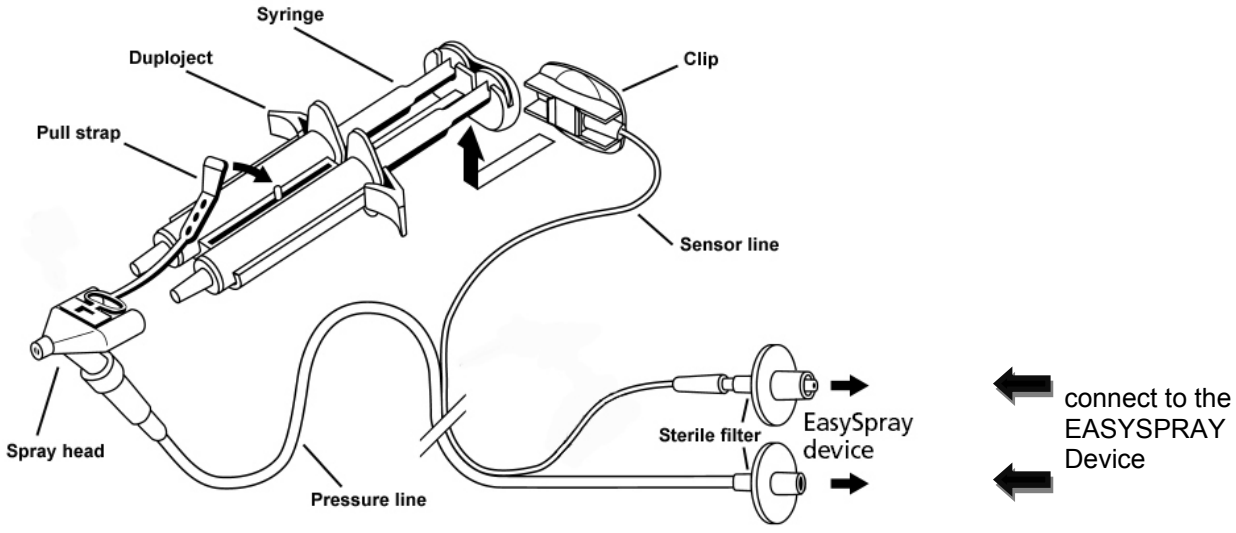
If application of the fibrin sealant components is interrupted, clogging may occur in the cannula. To resume application, replace the application cannula with a new one. If the apertures of the joining piece are clogged, use the spare joining piece provided in the Duo Set.

b) Simultaneous Application Using Spray Set and EASYSPRAY:

Caution must be used when applying fibrin sealant using pressurized air or gas.
- Any application of pressurized air or gas is associated with a potential risk of air embolism, tissue rupture, or gas entrapment with compression, which may be life-threatening.
- TISSEEL must be sprayed only onto application sites that are visible.

Spray Set and EASYSPRAY can be connected to TISSEEL Kit (see drawing below) and TISSEEL Frozen in the same way.

Note: For operation instructions please refer to the Instructions for Use provided together with Spray Set and Easy Spray.



The Spray Set is particularly suitable for spraying of larger areas, e.g. to control oozing of parenchymatous organs or to adhere skin grafts. The two components are sprayed simultaneously using sterile propellant gas via EASYSPRAY, and the volume of the Solutions ejected is controlled with the DUPLOJECT System / DUO Set plunger. The user must follow the instructions and precautions in the Spray Set and EASYSPRAY device user manual, for example regarding the need to limit the gas pressure to a maximum of 2 bars. Spray at a minimum distance of 10 cm. Only use application devices licensed for the administration of TISSEEL.

c) Simultaneous Application Using DUPLOCATH Application Catheters or other accessories provided by Baxter:

In operation sites where access is difficult or when using an endoscope or trocar, TISSEEL can be applied using DUPLOCATH Application Catheters.

Note: For operation instructions please refer to the Instructions for Use provided together with DUPLOCATH Application Catheter 25, 35 or 180.

In case of using other accessories provided by Baxter, please refer to the operating instructions contained in the Instructions for Use for the particular accessory.

d) Simultaneous Application by Premixing:

Mix equal volumes of the two components and immediately apply them to the recipient surface or surfaces. If desired, the Thrombin Solution can be mixed with spongiosa to pack bone defects. Hold in place for three minutes. Once turbid, TISSEEL can no longer be manipulated.

e) Simultaneous Application Using DuploSpray MIS Surgical Sealant Delivery System

The DuploSpray MIS applicator is intended for the application of TISSEEL Fibrin Sealant.

The applicator consists of a rigid and sterile stainless steel cannula, dual lumen spray tubing, sterile filter, and two replacement spray tips. It delivers a thin, uniform film of fibrin sealant over wound surfaces for maximum product utilization and broad area coverage.

The regulator is a pressure-regulating device for the control and release of compressed gas from a pressurized CO_2 source; this is the safest gas to use in laparoscopic procedures, and with a patient vent line, provides pressure stability during application. The gas-assisted applicator mixes and disperses the combined TISSEEL components for application of a thin layer to an open or

shallow treatment site. The spray output is adjustable from a coarse to fine droplet size and a foot switch activates the gas flow. When using the DuploSpray MIS spray device (for minimally invasive surgery) it should not be sprayed closer than 2 cm.

A diagram of the DUPLOSPRAY MIS APPLICATOR and DUPLOSPRAY MIS REGULATOR (original and updated models) attached to the DUPLOJECT double chamber syringe containing the two-component fibrin sealant is provided below:

1. Original DUPLOSPRAY MIS REGULATOR (x 1)
2. Stainless steel cannula (x 1)
3. Replaceable spray tip (x 2)
4. Dual lumen spray tubing and filter (x 1)
5. DUPLOJECT double chamber syringe (x 1) containing TISSEEL fibrin sealant

Figure 1: DUPLOSPRAY MIS APPLICATOR and Original DUPLOSPRAY MIS REGULATOR

Figure 2: Updated DUPLOSPRAY MIS REGULATOR. This model is available in Canada as of April 2012.

Note: For operation instructions please refer to the Instructions for Use provided together with DuploSpray MIS Applicator

Gluing of Tissue

After the two components have been applied, approximate the wound areas. Fix or hold the glued parts in the desired position for three to five minutes to ensure that the setting Sealant adheres firmly to the surrounding tissue. Solidified Sealant reaches its ultimate strength after about two hours (70% after about ten minutes).

1.9 OVERDOSAGE

No symptoms of overdose are known for TISSEEL. As the product is actively used by the surgeon, overdose is very unlikely to occur. TISSEEL is only to be used by a physician in a hospital setting.

Only a thin layer of TISSEEL should be applied to avoid the formation of excess granulation tissue and to ensure gradual absorption of the polymerized fibrin sealant. Excessive thickness of the fibrin layer may negatively interfere with the product's efficacy and the wound healing process (See *DOSAGE AND ADMINISTRATION* and *WARNINGS AND PRECAUTIONS*).

1.10 ACTION AND CLINICAL PHARMACOLOGY

Mechanism of Action

TISSEEL (Fibrin Sealant (Human), Vapor Heated, Solvent/Detergent Treated) is a tissue glue with sealing, hemostyptic and gluing properties, which does not interfere with but may enhance wound healing.

The action of TISSEEL simulates key features of the physiological process of wound closure. A highly concentrated fibrinogen Aprotinin solution, which among other ingredients contains Factor XIII co-fractionated from the plasma, and a solution of thrombin and calcium chloride are applied to the wound area, where the mixture coagulates. The presence of Factor XIII causes the fibrin to crosslink, which gives the coagulum additional resilience. Aprotinin prevents premature degradation of the clot.

Pharmacodynamics

Thrombin is a highly specific protease that transforms the fibrinogen contained in Sealer Protein (Human) into fibrin. Part of the thrombin is adsorbed by the fibrin, and any excess thrombin is inactivated by protease inhibitors in the blood.

Fibrinolysis Inhibitor, Aprotinin (Bovine or Synthetic), is a polyvalent protease inhibitor that prevents premature degradation of fibrin. Free Aprotinin has a half-life (t ½) of approx. 0.82 and is eliminated by the kidney. Preclinical studies with different fibrin sealant preparations simulating the fibrinolytic activity generated by extracorporeal circulation in patients during cardiovascular surgery have shown that incorporation of Aprotinin in the product formulation increases resistance of the fibrin sealant clot to degradation in a fibrinolytic environment.

Pharmacokinetics

TISSEEL is intended for local application only, therefore systemic exposure or distribution to other organs or tissues is not expected and pharmacokinetic studies were not conducted.

1.11 STORAGE AND STABILITY

TISSEEL Kit (Freeze-Dried)

When stored between 2°C and 25°C (36°F and 77°F), TISSEEL (Fibrin Sealant (Human), Vapor Heated, Solvent/Detergent Treated) Kit, lyophilized, is stable until the expiry date indicated on the label. Do not freeze.

After reconstitution, the product must be used within 4 hours. Reconstituted solutions must not be refrigerated or frozen.

TISSEEL Pre-filled Syringe (Frozen)

When stored at ≤ -20°C (≤ -4°F), TISSEEL (Fibrin Sealant (Human), Vapor Heated, Solvent/Detergent Treated) Frozen, in pre-filled syringes, is stable until the expiry date indicated on the label.

Room Temperature Storage: Unopened pouches, thawed at room temperature, may be stored for up to 7 days at room temperature (15°C -25°C) after removal from the freezer.

1.12 DOSAGE FORMS, COMPOSITION AND PACKAGING

Dosage Forms and Packaging

TISSEEL (Fibrin Sealant (Human), Vapor Heated, Solvent/Detergent Treated) is available in two different dosage forms: Kit and Frozen.

Strength	Volume (Total)	Kit	Frozen
500 IU/mL	2 mL 4 mL 10 mL	One vial lyophilized Sealer Protein (Human) Concentrate, to be reconstituted with one vial 1.0 mL/ 2.0 mL/ 5.0 ml of Aprotinin Solution, sterile One vial lyophilized Thrombin (Human) 500 IU, to be reconstituted with one vial 1.0 mL/ 2.0 mL/ 5.0 mL of Calcium Chloride Solution, sterile	1.0 mL/ 2.0 mL/ 5.0 mL of sterile frozen TISSEEL Sealer Protein (Human)-Aprotinin Solution 1.0 mL/ 2.0 mL/ 5.0 mL of sterile frozen Thrombin (Human) – Calcium Chloride Solution

TISSEEL Kit also contains DUPLOJECT System, the Kit for Reconstitution and Application consisting of 4 transfer needles, 2 blue-scaled syringes, 2 black-scaled syringes, 1 DUPLOJECT two-syringe clip, 2 joining pieces and 4 application cannulas.

TISSEEL Frozen also contains DUO Set, the sterile accessory devices consisting of 1 plunger, 2 joining pieces and 4 application cannulas.

See the following *Accessories* Section for more accessories for use with TISSEEL .

Composition

TISSEEL 500 IU Kit consists of 4 separate vials containing Sealer Protein (Human), Aprotinin (bovine or synthetic) Solution, Thrombin (Human) and Calcium Chloride Solution.

TISSEEL 500 IU Frozen consists of a double-chamber syringe containing Sealer Protein-Aprotinin (bovine or synthetic) Solution (syringe body 1) and Thrombin-Calcium Chloride Solution (syringe body 2).

After reconstituting the vials in the Kit, the resulting concentrations of the Sealer Protein-Aprotinin solution and the Thrombin-Calcium Chloride solution are the same as those in the Frozen presentation.

	Component	Amount
1	Sealer Protein (Human)-Aprotinin Solution, sterile, contains:	
	- Total protein	96-125 mg/mL
	- Factor XIII	0.6-10 U/mL**
	- Fibrinogen (Clottable Protein)	72-110 mg/mL
	- Plasmafibronectin (CIG)*	2-9 mg/mL
	- Plasminogen*	40 – 120 µg/mL
2	Aprotinin (bovine or synthetic) Solution, sterile	3,000 KIU/mL***
3	Thrombin (Human)	500 IU/mL ****
4	Calcium Chloride Solution, sterile	40 µmol/mL

* Development data, not tested at the Final Product level
** One unit corresponds to the amount of Factor XIII contained in 1 mL of fresh normal plasma.
*** 30 Kallidinogenase Inactivator Units (KIU) correspond to 1 FIP-Unit[18].
**** One International Unit (IU) of Thrombin is defined as the activity contained in 0.0853 mg of the First International Standard of Human Thrombin[19].

Nonmedicinal ingredients in the Sealer Protein-Aprotinin solution are human albumin, niacinamide and water for injection (WFI). Nonmedicinal ingredients in the Thrombin-Calcium Chloride solution are human albumin, sodium chloride and water for injection (WFI). Sodium Hydroxide and Hydrochloric Acid are used to adjust the pH in both solutions.

All plasma units used for manufacture are ALT tested and non-reactive in tests for Hbs-antigen and antibodies to HCV, HIV-1 and HIV-2. Before further processing all individual plasma donations are subjected to an inventory hold for a possible look-back of plasma donations suspected of infection. All plasma units are tested by HIQ-PCR = Hyland Immuno Quality Assured Polymerase Chain Reaction.

1.13 ACCESSORIES

The following are some accessories for use with TISSEEL VH S/D. A complete list of accessories can be obtained from a Baxter representative. When using these devices, strictly follow the Instructions for Use of the devices.

FIBRINOTHERM	Combined Heating and Stirring Device
EASYSPRAY	Propellant gas control unit, manometer, reducing valve, and pressure tube.
Spray Set (sterile, disposable)	Disposable set consisting of sterile filter with connection tube, sensing line and a spray head.
DUPLOCATH 25 Application Catheter	Length: approximately 25 cm (10”) Diameter: approximately 5 french (approx. 0.17 cm) Radiopaque. Sterile. Disposable.
DUPLOCATH 35 M.I.S Application Catheter	Catheter: Length: approximately 35 cm (14”) Diameter: approximately 5 french (approx. 0.17 cm) Adapter: Length: approximately 30 cm (12”) Diameter: 15 french (0.5 cm) For insertion through a 5-6 mm trocar in minimally invasive surgery (M.I.S.) Radiopaque. Sterile. Disposable.
DUPLOCATH 180 Application Catheter	Length: approximately 180 cm (70”) Diameter: approximately 5 french (approx. 0.17 cm) For use with an endoscope. The Application Catheter 180 can be shortened to any length necessary. Radiopaque. Sterile. Disposable.
DUPLOSPRAY MIS SURGICAL SEALANT DELIVERY SYSTEM	Propellant gas control unit with applicators and tips for the application of surgical sealant.
DuploTip Dual Lumen Cannula	Catheter double canal Length: 10.4 cm or 32 cm

Alternative reconstitution accessories are available for TISSEEL.

2. PART II: SCIENTIFIC INFORMATION

2.1 PHARMACEUTICAL INFORMATION

Drug Substance
See PART I – DESCRIPTION and PART II – DETAILED PHARMACOLOGY Sections.

Product Characteristics
See PART I – DESCRIPTION and PART II – DETAILED PHARMACOLOGY Sections.

Viral Inactivation

TISSEEL (Fibrin Sealant (Human), Vapor Heated, Solvent/Detergent Treated) is made from human plasma. Products made from human plasma may contain infectious agents, such as viruses, that can cause disease. The risk that such products will transmit an infectious agent has been reduced by screening plasma donors for prior exposure to certain viruses (see Detailed Pharmacology), by testing for the presence of certain current virus infections, and by inactivating and removing certain viruses in the course of the manufacture.

The manufacturing procedure for TISSEEL includes processing steps designed to further reduce the risk of viral transmission. In particular, vapor heating and solvent/detergent treatment processes are included in the manufacturing of Sealer Protein Concentrate and Thrombin. In addition the reduction factors as associated with DEAE-Sephadex Batch Chromatography and Ion Exchange Chromatography of the Thrombin drug substance purification were investigated. Validation studies were conducted using samples drawn from manufacturing intermediates for each of the two human plasma derived components. These samples were spiked with stock virus suspensions of known titers followed by further processing under conditions equivalent to those in the respective manufacturing steps.

The virus reduction factors (expressed as $\log_{10}$) of independent manufacturing steps were as follows for each of the viruses tested:

Reduction Factors for Virus Removal and/or Inactivation Sealer Protein Component						
	Mean Reduction Factors [log_{10}] of Virus Tested*					
Manufacturing Step	HIV-1	HAV	BVDV	PRV	MMV	B19V
Early Manufacturing Steps	n.d.	n.d.	n.d.	n.d.	2.7**	3.4**
					2.3**	
Solvent/Detergent Treatment	>5.3	n.d.	>5.7	>5.9	n.d.	n.d.
Vapor Heat Treatment	>5.5	>5.6	>5.7	>6.7	1.2	1.0
Overall Reduction Factor (ORF)	**>10.8**	**>5.6**	**>11.4**	**>12.6**	**3.5**	**3.3**
Reduction Factors for Virus Removal and/or Inactivation Thrombin Component						
	Mean Reduction Factors [log_{10}] of Virus Tested*					
Manufacturing Step	HIV-1	HAV	BVDV	PRV	MMV	B19V
Removal of Thrombin precursor protein from Cryosupernatant	3.2	1.5	1.8	2.5	1.2	1.7
Vapor Heat Treatment	>5.5	>4.9	>5.3	>6.7	1.0	>4
Solvent/Detergent Treatment	>5.3	n.d.	>5.5	>6.4	n.d.	n.d.
Ion Exchange Chromatography	n.d.	n.d.	n.d.	n.d.	3.6	n.d.
Overall Reduction Factor (ORF)	**>14.0**	**>6.4**	**>12.6**	**>15.6**	**5.8**	**>5.7**

n.d. = not determined

* The mean RF of all runs was calculated. E.g. when two runs were performed in study A and one run in study B for HIV-1, then the mean of the three RF's was calculated and listed in the table.

** As a conservative value for general and robust Parvovirus reduction capacity in steps 1 to 8, 2.3 logs were calculated from MMV and B19V reduction factors (omitting the higher B19V reduction factor). For calculation of and rationale behind this value, see further above in the section where the corresponding study is discussed. For calculation of virus-specific overall reduction factors, however, only virus-specific individual reduction factors were considered.

HIV-1: Human immunodeficiency virus 1; **HAV**: Hepatitis A virus; **BVDV**: Bovine viral diarrhea virus, a model for Hepatitis C virus; **PRV**: Pseudorabies virus, a model for enveloped DNA viruses, among those Hepatitis B virus; **MMV**: Mice minute virus, a model for Human Parvovirus B19, **B19V**: Human Parvovirus B19.

2.2 CLINICAL TRIALS

Numerous clinical studies investigating the safety and efficacy of the product as a hemostyptic and biodegradable tissue glue in various fields of surgery have been performed. A number of these were controlled studies in fields including orthopedic surgery, abdominal surgery, urology, and cardiovascular surgery. The cardiovascular safety study using the heat treated product has shown that TISSEEL (Fibrin Sealant (Human), Vapor Heated, Solvent/Detergent Treated) transmits neither hepatitis viruses nor HIV. Pre-clinical studies have shown that the vapor treated product is at least as effective as the heat treated product. In the clinical study 550003 described below, TISSEEL was shown to be non-inferior to an earlier formulation of the product, TISSEEL VH (predecessor product) and both products were shown not to transmit hepatitis, HIV or P19B.

Use of TISSEEL has invariably shown superior results in the groups treated as against the untreated controls who underwent the same types of surgery. These results were attributable to an improved hemostasis and, therefore, reduced blood loss, a tighter sealing of sutures preventing leakages or a fast and uncomplicated healing of the surgical wound.

In none of the studies have systemic side-effects been seen nor has any product related transmission of viral hepatitis or HIV occurred in any of the patients treated.

TISSEEL 500 IU (Fast Set):

TISSEEL 500 IU was evaluated in a prospective, parallel design, randomized (1:1), double-blind, multicenter clinical study (550003) against an earlier formulation of the product, TISSEEL VH, in 317 subjects undergoing cardiac surgery requiring cardiopulmonary bypass (CPB) and median sternotomy. Patients were treated with TISSEEL 500 IU or the control product only when hemostasis was not achieved by conventional surgical methods. For the endpoint, hemostasis achieved at the primary treatment site within 5 minutes of treatment and maintained until closure of the surgical wound, TISSEEL 500 IU was at least as effective as the earlier formulation of the product using a one-sided 97.5% confidence interval on the difference in the proportion of subjects successfully treated.

Hemostasis within 5 minutes and maintained until surgical closure		
	TISSEEL 500 IU	**Predecessor product TISSEEL VH**
Intent to Treat Analysis	127/144 (88.2%)	129/144 (89.6%)
Per Protocol Analysis	108/123 (87.8%)	122/135 (90.4%)

Geriatrics (> 65 years of age):

In this Phase 3 clinical study of TISSEEL 500 IU, 71 of 144 subjects were 65 and over. No overall differences in safety or effectiveness were observed between these subjects and younger subjects, and other reported clinical experience has not identified differences in responses between the elderly and younger patients.

An earlier formulation of TISSEEL's predecessor product, TISSEEL HT (heat treated, containing bovine thrombin) was evaluated in an open-label crossover study against control topical hemostatic agents in 489 patients undergoing cardiovascular reoperation or resternotomy at 11 institutions. Patients were randomized to TISSEEL HT or control hemostatic agents when a topical hemostatic was needed at the conclusion of surgery and after all attempts at surgical hemostasis. Patients were crossed to the alternative therapy if bleeding continued after the 5 minute endpoint. At 10 centres, TISSEEL HT was used after administration of protamine sulfate. At one site, TISSEEL HT could be used before administration of protamine sulfate. 365 of the 489 patients developed bleeding episodes requiring treatment. In these patients, for the endpoint, successful hemostasis at 5 minutes, TISSEEL HT was statistically significantly superior to control topical hemostatic agents.

Hemostasis within 5 minutes	
Predecessor TISSEEL HT	**Control Topical Hemostatic Agent**
82.4% (159/193)	44.5% (76/172)
Pearson χ^2 two sided; p <0.0001; intent-to-treat analysis	

Similarly, absolute time to cessation of bleeding was statistically significantly shorter for TISSEEL HT than for control topical hemostatic agents (p<0.0001, Wilcoxon-Gehan test, two sided).

In a single center, open label trial, an earlier formulation of TISSEEL HT was compared to historical controls in patients undergoing laparotomy for blunt or penetrating traumatic injury to the spleen and/or liver. Use of TISSEEL HT resulted in the need for statistically significantly fewer splenectomies than control hemostatic maneuvers:

Splenectomy Rate			
Injury to:	**Predecessor TISSEEL HT**	**Historic Controls**	
Spleen	0/19	14/22	p <0.001
Spleen and liver	1/26	19/34	p <0.001

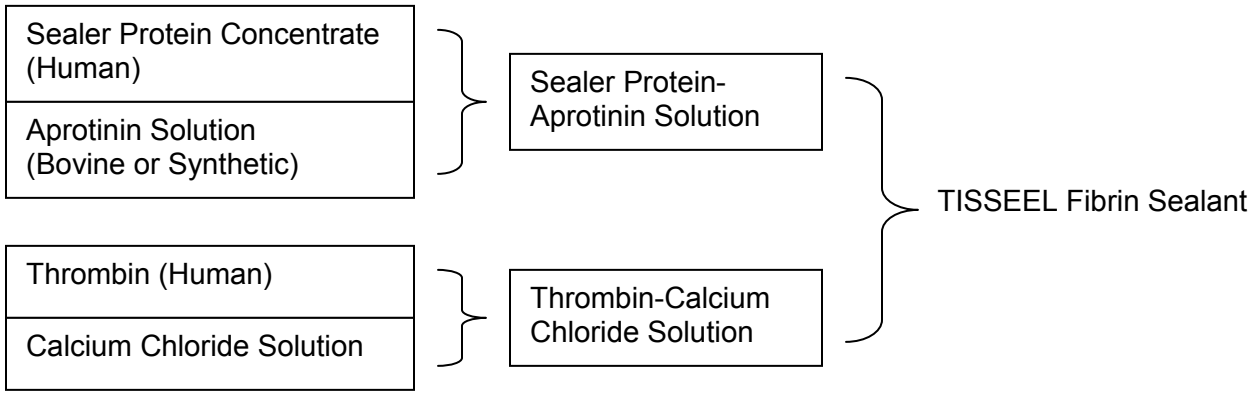
TISSEEL HT did not result in significantly reduced mortality in patients with blunt or penetrating trauma to the liver alone or to the liver and spleen (p=0.067, χ^2, one sided).
In a single center, prospective open label study of 120 patients randomized to standard of care (59 patients) or standard of care plus fibrin sealant (61 patients) for elective colostomy closure after temporary colostomy placement for treatment of traumatic injury to the colon, the earlier

version of TISSEEL HT plus standard of care was also shown to be significantly superior to standard of care alone (p=0.0406, Jonckheere-Terpstra test for ordinal data, two sided) with regard to anastomotic complications (leakage, intra-abdominal abscess formation, re-operation, septic shock, and death).

2.3 DETAILED PHARMACOLOGY

The substances in the TISSEEL (Fibrin Sealant (Human), Vapor Heated, Solvent/Detergent Treated) Kit are used to prepare two components: Sealer Protein Solution and Thrombin Solution. Sealer Protein Solution is produced by dissolving Sealer Protein Concentrate in Aprotinin Solution. Freeze-dried Thrombin, dissolved in Calcium Chloride Solution, yields the Thrombin Solution.

The assembly for TISSEEL Frozen is reflecting the composition of the Kit components after reconstitution, filled in a double-chamber syringe, which is stored in a deep frozen condition: Sealer Protein Concentrate-Aprotinin (syringe chamber 1) and Thrombin-Calcium Chloride Solution (syringe chamber 2).



The two components are mixed either immediately before application to the recipient surface or in situ using one of the methods described under Method of Application. The Sealer Protein / Thrombin Solution, after mixing, is a viscous solution adhering to wound surfaces and quickly sets to form a white, rubberlike mass, which continues to gain in strength within two hours following application. This process is made use of to achieve hemostasis, and to seal or glue tissue.

In this manner, the need for sutures may be reduced, although not totally eliminated. The time until Sealant sets can be used to approximate wound edges, to provide optimum conditions for healing. In the course of wound healing, Sealant is completely absorbed.

Fibrin Sealant is a two-component biological sealant produced from pooled human plasma. As for all plasma products, the following measures are implemented to ensure the safety of the product from the potential presence of pathogenic viruses in human plasma:

- Donor selection
- Donation testing of single donations and also at the mini-pool and manufacturing pool level
- The use of PCR assay system (PCR) for release of plasma pools
- Effective virus inactivation/removal steps integrated into the manufacturing process including validation

Each plasma donation is tested for infectious markers for human immunodeficiency virus, types 1 and 2 (HIV-1/-2), hepatitis C virus (HCV) and hepatitis B surface antigen (HbsAg) The criteria for release of each single plasma donation for further manufacturing are as follows:

- HIV-1 / HIV-2 antibody: non-reactive
- HbsAg: non-reactive
- HCV antibody: non-reactive

Each manufacturing plasma pool is tested and released for further manufacturing only when

- non-reactive for Human Immunodeficiency Virus (HIV), Hepatitis C Virus (HCV), Hepatitis A Virus (HAV) and Hepatitis B Virus (HBV) using Nucleic Acid Testing (NAT)
- Parvovirus B19 concentration not exceeding 10^4 IU / mL as measured by NAT

Plasma pool are tested using HIQ-PCR = Hyland Immuno Quality Assured Polymerase Chain Reaction. With the PCR method, in general 500 genome equivalents/mL of the above viruses can be determined reliably, with the actual sensitivity of HIQ-PCR being below that. Therefore all pools which have been tested and evaluated as being positive lead to exclusion from further processing.

The plasma safety precautions and virus inactivation/removal steps taken during the manufacture of TISSEEL VH S/D 500 IU are equivalent to those used for ARTISS 4 IU. During the pivotal clinical study of TISSEEL VH S/D 500 IU Fibrin Sealant there were zero confirmed seroconversions for HAV, HBV, HCV, HIV or B19V (Baxter clinical study 550003).

Although viral safety was not an endpoint in the other clinical studies, any issues concerning viral safety would have been detected and followed up in the process of AE reporting. No reports concerning viral transmission from any studies, involving Fibrin Sealant 4 IU (ARTISS) or 500 IU (TISSEEL) were received.

The therapeutic activities of TISSEEL are hemostasis, gluing and sealing of tissue, and the support of wound healing.

Physiologically, the process of wound closure starts when a bleeding ceases. In places where injured blood vessels lie open, hemostatic plugs form of platelets and fibrin, which become more and more solid as other blood cells, particularly erythrocytes, are involved. Because the bleeding ceases, blood that has escaped into the wound bed earlier coagulates. In a next step fibrin in both the coagulated blood and hemostatic plug retracts, plasma is squeezed out, the blood vessels contract, and the wound area becomes smaller. As various cells begin to proliferate into the retracted blood coagula, wound healing sets in.

In developing Fibrin Sealant, this principle has at least partly been simulated by applying a highly concentrated fibrinogen solution, which also contains factor XIII as co-fractionated from the plasma, and a solution of thrombin and calcium chloride to the wound area, where the mix coagulates. Since the Sealant does not contain thrombocytes, the clotted fibrin does not appreciably retract. When the mixture is applied, however, the fibrin concentration in the coagulum is the same as or higher than that of a retracted hemostatic plug. To preserve the coagulum until the wound healing has reached a stage where it is no longer required, the Sealant has been designed to contain a fibrinolysis inhibitor.

The fibrin in the Sealant adheres perfectly to the wound edges guaranteeing adequate sealing effect. Similar to the hemostatic plug, wound healing is promoted by the fibrin applied. Combined application of Sealant with a mixture of autologous or homologous spongiosa provides excellent plugging material for bone defects. Using adequate technique, TISSEEL is also an excellent tool in sealing autologous or homologous cartilage and bone.

TISSEEL and all of its components have not been observed to affect systemic circulation, respiration, or the central nervous system. This is attributable to both, the fact that only minimal quantities of each single component are applied compared to their use for other indications, and the fact that TISSEEL becomes only locally effective.

The mechanism underlying solidification of tissue and persistence of a solidified clot have been investigated in numerous studies.

As a biologic material, Fibrin Sealant becomes completely absorbed at a rate which depends both on the fibrinolytic activity of the surrounding tissue and the quantity of Fibrinolysis Inhibitor added. In the course of wound healing the Sealant clot is gradually replaced by ingrowing tissue, Thrombin is inactivated by the physiological protease inhibitors, Calcium Chloride is subjected to the calcium and chloride catabolism of the organism, and Aprotinin and its metabolites are eliminated by the kidney.

2.4 TOXICOLOGY

The local application of TISSEEL (Fibrin Sealant (Human), Vapor Heated, Solvent/Detergent Treated) underlines the importance of histological studies for toxicology data. Accordingly, histologies have been performed on various species in tissues ranging from skin, vessels, nerves, tendons, organ tissue to bone.

In vivo toxicology studies indicated no acute toxicity and normal local tolerance reactions of TISSEEL Frozen[2] with bovine or synthetic Aprotinin in rats and rabbits, respectively. Furthermore, no difference between synthetic and bovine Aprotinin for single-dose toxicity after intravenous application could be detected in mice and rats and synthetic Aprotinin was well tolerated in rabbits.

No skin sensitizing potential of synthetic Aprotinin was shown in a Guinea Pig Maximization Test.

In vitro, no differences were found between TISSEEL 500 IU Frozen with bovine or synthetic Aprotinin and the predecessor product TISSEEL VH 500 IU lyophilized Kit with respect to cellular compatibility, or mutagenicity.

Synthetic Aprotinin was non-mutagenic an Ames test.

Cellular compatibility was confirmed in an additional *in vitro* study comparing both TISSEEL preparations (frozen and lyophilized) and the predecessor product TISSEEL VH (lyophilized) Kit.

In summary, presented *in vitro* and *in vivo* data support the equivalence of TISSEEL (frozen and lyophilized) with the predecessor product TISSEEL VH (lyophilized Kit). Moreover, it was shown that synthetic Aprotinin has no negative impact on the safety of the Fibrin Sealant.

Long-term animal studies to evaluate the carcinogenic potential of TISSEEL or studies to determine the effect of TISSEEL on fertility have not been performed.

On the following page the toxicology program is presented.

[2] The composition of TISSEEL frozen, thawed, is identical to the composition of the reconstituted TISSEEL freeze-dried.

Toxicology program for Baxter Fibrin Sealants

Study Type	Route of Administration	Species	Test Article
Single-dose Toxicity	s.c.	Rat	TISSEEL (frozen)
Single-dose Toxicity	s.c.	Rabbit	TISSEEL (frozen)
Single-dose Toxicity	s.c.	Rat	TISSEEL s-apr (frozen)
Single-dose Toxicity	s.c.	Rabbit	TISSEEL s-apr (frozen)
Single-dose Toxicity	i.v.	Mouse	Synthetic Aprotinin
Single-dose Toxicity	i.v.	Rat	Synthetic Aprotinin
Genotoxicity	i.v.	*E.coli*	TISSEEL (frozen)
Genotoxicity	i.v.	*Salmonella typhimurium*	Synthetic Aprotinin
Local Tolerance	s.c. spongiosa blocks	Rat	TISSEEL (frozen)*
Local Tolerance	s.c. spongiosa blocks	Rabbit	TISSEEL (frozen)*
Local Tolerance	s.c. spongiosa blocks	Rat	TISSEEL s-apr (frozen)*
Local Tolerance	s.c. spongiosa blocks	Rabbit	TISSEEL s-apr (frozen)*
Local Tolerance	i.v., paravenous	Rabbit	Synthetic Aprotinin
Cytotoxicity	i.v.	Human lung fibroblasts	TISSEEL (frozen)
Cytotoxicity	i.v.	Human lung fibroblasts	TISSEEL (lyophilized)
Cytotoxicity	i.v.	Human lung fibroblasts	TISSEEL s-apr (frozen)
Skin Sensitization	intradermal, epicutan	Guinea pig	Synthetic Aprotinin

* Local tolerance studies using a subcutaneously implanted spongiosa block model were conducted with fibrin sealants diluted to a Thrombin concentration of 4 IU/mL.

3. PART III: CONSUMER INFORMATION

3.1 PATIENT COUNSELLING INFORMATION

Because this product is made from human plasma, the physician should discuss the risks and benefits with the patient. Patient should be encouraged to consult with their physician if symptoms of B19 virus infection appear (fever, drowsiness, chills and runny nose followed about two weeks later by a rash and joint pain).

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